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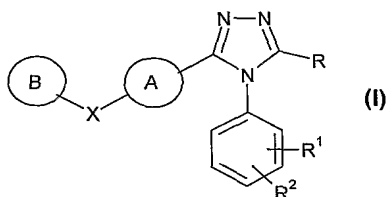
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(54) Title: 3-HETEROCYCLYL-4-PHENYL-TRIAZOLE DERIVATIVES AS INHIBITORS OF THE VASOPRESSIN V1A RE-
CEPTOR



(57) Abstract: Compounds of formula (I): or a pharmaceutically acceptable
derivative thereof, wherein R represents C₁₋₆alkyl (optionally substituted
by C₁₋₆alkyloxy or Het) or C₁₋₆alkyloxy; R¹ and R² independently represent
hydrogen, halo or C₁₋₆alkyl, ring A represents Het¹; X represents O or NR³; R³
represents hydrogen or C₁₋₆alkyl; ring B represents a phenyl group or Het², either
of which may be optionally substituted with one or more groups selected from
halo, CN, C₁₋₆alkyloxy, CF₃, C₁₋₆alkyl, NH₂ and NO₂; Het and Het¹ independently
represent a 5- or 6-membered saturated, partially unsaturated or aromatic
heterocyclic group comprising either (a) 1 to 4 nitrogen atoms, (b) one oxygen or

one sulphur atom or (c) 1 oxygen atom or 1 sulphur atoms and 1 or 2 nitrogen atoms are useful for treating anxiety, cardiovascular
disease (including angina, atherosclerosis, hypertension, heart failure, edema, hypernatremia), dysmenorrhoea (primary and
secondary), endometriosis, emesis (including motion sickness), intrauterine growth retardation, inflammation (including rheumatoid
arthritis) mittlemerchz, preclampsia, premature ejaculation, premature (preterm) labour and Raynaud's disease.